(12) UK Patent Application (19) GB (11)

2039938 A

- (21) Application No 7931601
- (22) Date of filing 12 Sep 1979
- (30) Priority data
- (31) 7826343
- (32) 13 Sep 1978
- (33) France (FR)
- (43) Application published 20 Aug 1980
- (51) INT CL³ C11D 3/37
- (52) Domestic classification C5D 6A5C 6B12N1 6B12N2 6B12N5 6B12NX 6B12P 6B13 6B1 6C8
- (56) Documents cited GB 1437912 GB 1228060
- (58) Field of search C5D
- (71) Applicants L'Oreal, 14 rue Royale, 75008 Paris, France.
- (72) Inventors
 Guy Vanlerberghe,
 Henri Sebag,
 Alexandre Zysman,
 Claude Dubief.
- (74) Agents J.A. Kemp & Co.

- (54) Composition for treating fibrous materials, based on cationic and anionic polymers
- (57) Compositions for the treatment of fibrous materials contain at least one cationic polymer, at least one anionic polymer, at least one alkali metal salt and at least one non-ionic surfaceactive agent or surface-active agent which contains one or more carboxyl or carboxylate groups in addition to non-ionic groups.

20

35

SPECIFICATION

Composition for treating fibrous materials, based on cationic and anionic polymers

5 The present invention relates to compositions based on polymers, which are intended to be used in the treatment of fibres, in particular hair or textile fibres. It relates more particularly to compositions for washing or dyeing these materials.

5

Numerous compositions, in particular cosmetic compositions, containing either anionic polymers or cationic polymers, are known, the purpose of which is to modify the properties of the materials treated.

We have discovered that it is very advantageous to treat fibrous materials with compositions which simultaneously contain one or more anionic polymers, one or more cationic polymers, alkali metal salts and one or more non-ionic surface-active agents or surface-active agents which contain one or more carboxyl or carboxylate groups in addition to non-ionic groups i.e. a non-ionic surface-active molecule which is substituted by one or more carboxyl or carboxylate groups.

10

In fact, by using this combination, it is possible to impart, for example to hair, softness and ease of 15 comb-out at the same time as waveset hold, strength or gloss. Textile fibres treated with this combination also possess valuable properties of softness and good hold.

15

These compositions exhibit the advantage of being homogeneous and stable; when they are diluted with water on rinsing the hair or textile materials, they produce a deposit of polymers on these materials.

In certain cases, the problem of the solubilisation of the precipitate which can form can be solved by the use of an appropriate solubilising agent, such as an organic solvent or anionic surface-active agent, and by suitably adjusting the pH. We have discovered that, by using, according to the present invention, alkali metal salts and non-ionic surface-active agents or surface-active agents which contain one or more carboxyl or carboxylate groups in addition to non-ionic groups, it is possible to obtain compositions which are homogeneous and stable at a pH which is lower than that required when using only the above-mentioned

20

solubilising agents, and is substantially closer to neutrality, preferably from 5 to 8, which compositions are capable of producing a deposit of polymers on dilution with water.

25

The present invention therefore provides a composition which is suitable for use, and intended to be used, in the treatment of hair and textile fibres and which contains at least one anionic polymer, at least one 30 cationic polymer, at least one alkali metal salt and at least one non-ionic surface-active agent or surface-active agent which contains one or more carboxyl or carboxylate groups in addition to non-ionic groups.

30

The present invention also provides a process for treating hair or textile materials, which employs a composition of this invention.

The composition of the present invention generally has a pH of 5 to 8.

35

The cationic and anionic polymers are preferably each present in an amount of 0.25 to 3% by weight, the alkali metal salt is present in an amount of 0.25 to 8% by weight and the said surface-active agent is present in an amount of 1 to 50% by weight, preferably 5 to 25% by weight.

The ratios of the said cationic polymer to the anionic polymer, present in the compositions according to 40 the invention, are preferably from 5 to 0.4, in particular from 3 to 0.5, expressed as the ratio of equivalents of cationic units to equivalents of anionic units.

The alkali metal salts which are particularly preferred according to the invention are sodium, potassium or lithium salts. These salts are preferably chosen from amongst halides, such as chloride and bromide, sulphates, or salts of organic acids, such as acetates or lactates.

The anionic polymers which are particularly preferred according to the invention are polymers which must 45 be soluble in water after neutralisation of the acid groups with an alkali, such as sodium hydroxide or potassium hydroxide, or an amine, such as triethanolamine, 2-amino-2-methylpropan-1-ol or 2-amino-2methylpropane-1,3-diol, in the presence of the above-mentioned alkali metal salts and surface-active agents, in proportions by weight of, for example 1/0.25 to 8/3 to 30, respectively.

Anionic polymers which can be used according to the invention include polymers which contain several carboxylic acid groups in their chain.

55

The carboxylic acid groups are generally provided by unsaturated mono- or di-carboxylic acids, as monomers, such as those corresponding to the formula

55

$$R_1$$
 C COOH
$$R_3$$
 (1)

in which n is 0 or an integer from 1 to 10, A denotes a methylene group joined to the carbon atom of the unsaturated group and/or to the adjacent methylene group if n is greater than 1, either directly or via a heteroatom, such as oxygen or sulphur, R1 denotes a hydrogen atom or a phenyl or benzyl group, R2 denotes a hydrogen atom or a lower alkyl or carboxyl group and R3 denotes a hydrogen atom, a lower alkyl group, a CH₂-COOH group or a phenyl or benzyl group.

In the above-mentioned formula, a lower alkyl radical preferably denotes a group having 1 to 4 carbon atoms, in particular methyl or ethyl.

65 The preferred anionic polymers according to the invention are:

35

40

45

homopolymers or copolymers of acrylic or methacrylic acid, in particular the products sold under the names VERSICOL E or K by ALLIED COLLOID or the name ULTRAHOLD 8 by CIBA GEIGY; the copolymers of acrylic acid and acrylamide sold in the form of their sodium salt under the names RETEN 421, 423 or 425 by HERCULES; and the acrylic or methacrylic acid - vinyl alcohol copolymers sold under the name HYDAGEN F by HENKEL;

copolymers of the above-mentioned acids with an unsaturated monoethylenic monomer, such as ethylene, vinylbenzene, vinyl and allyl esters and acrylic or methacrylic acid esters, which copolymers are optionally grafted to a polyalkylene glycol, such as polyethylene glycol, and optionally crosslinked. Such polymers are described, in particular, in French Patent No. 1,222,944 and German Application No. 2,330,956; 10 copolymers of this type containing, in their chain, an optionally N-alkylated and/or -hyroxyalkylated acrylamide unit, such as described, in particular, in Luxembourg Patent Applications Nos. 75,370 and 75,371, or sold under the name QUADRAMER 5 by American Cyanamid;

copolymers derived from crotonic acid, such as those containing, in their chain, vinyl acetate or propionate units and optionally units from other monomers, such as an allyl or methallyl ester, a vinyl ether or a vinyl ester of a saturated carboxylic acid having a long hydrocarbon chain, such as those containing at least 5 carbon atoms, it being possible for these polymers to be optionally grafted and crosslinked. Such polymers are described, inter alia, in French Patents Nos. 1,222,944, 1,580,545, 2,265,782, 2,265,781 and 1,564,110. Commercial products belonging to this class are the resins 28-29-30 and 26-13-14 sold by the Société National Starch; and

polymers derived from maleic, fumaric and itaconic acid or anhydride with vinyl esters, vinyl ethers, vinyl halides, phenylvinyl derivatives and acrylic acid and its esters, which polymers are described, in particular, in United States Patents Nos. 2,047,398, 2,723,248 and 2,102,113 and British Patent Specification No. 837,805, and especially those sold under the names GANTREZ AN or ES by General Anilin or the name EMA 1325 by MONSANTO; polymers which also belong to this class are the copolymers, described in French Patent Applications Nos. 76/13,929 and 76/20,917, of maleic, citraconic or itaconic anhydride and an allyl or methallyl ester and, optionally, acrylamide or methacrylamide which is monoesterified or monoamidated.

The cationic polymers are polymers of the polyamine or quaternary polyammonium type, the amine or ammonium group either forming part of the polymer chain or being joined to the latter.

Polymers of this type which can be used according to the invention include, vinylpyrrolidone/ 30 aminoalcohol acrylate copolymers (which may or may not be quaternised), such as those sold under the name GAFQUAT, for example "GAFQUAT 734 or 755", by the GAF Corp., which are described in greater detail, in particular, in French Patent No. 2,077,143, cellulose ether derivatives containing quaternary ammonium groups, such as those described in French Patent No. 1,492,597, and, in particular, the polymers sold under the name JR, such as JR-125, JR-400 and JR-30M, by UNION CARBIDE CORPORATION. Cationic polymers which give particularly valuable results include:

water-soluble cyclic polymers having a molecular weight of 20,000 to 3,000,000, such as polymers containing units corresponding to the formula (II) or (II');

$$(II) = \begin{bmatrix} CH_2 & -R^*C & CH_2 \\ H_2C & CH_2 \end{bmatrix} CH_2 + R^*C & CH_2 \\ R & R & CH_2 \end{bmatrix}$$

in which R" denotes hydrogen or methyl, R and R' independently of one another denote an alkyl group having from 1 to 22 carbon atoms, a hydroxyalkyl group in which the alkyl group preferably has 1 to 5 carbon atoms, or a lower amidoalkyl group, or R and R' together denote, together with the nitrogen atom to which they are attached, a heterocyclic group such as piperidinyl or morpholinyl, and Y is an anion such as bromide, chloride, acetate, borate, citrate, tartrate, bisulphate, bisulphate, sulphate or phosphate.

Amongst the quaternary ammonium polymers of the type defined above, those which are more particularly preferred are the dimethyldiallylammonium chloride homopolymers having a molecular weight of less than 100,000 which are sold under the name MERQUAT 100, and the copolymers of dimethyldiallylammonium chloride and acrylamide, having a molecular weight of more than 500,000 which are sold under 55 the name MERQUAT 550 by MERCK.

These cyclic polymers are described in French Patent No. 2,080,759 and its Certificate of Addition No. 2,190,406.

(2) homopolymers or copolymers derived from acrylic or methacrylic acid and containing the unit:

15

20

25

30

35

40

45

5

15

20

25

35

40

45

50

55

10 in which R₁ is H or CH₃, A is a linear or branched alkyl group having 1 to 6 carbon atoms or a hydroxyalkyl group having 1 to 4 carbon atoms, R₂, R₃ and R₄, which are identical or different, denote alkyl groups having 1 to 18 carbon atoms or a benzyl group, R₅ and R₆ are H or alkyl having 1 to 6 carbon atoms, and X⁻ denotes a methosulphate anion or a halide anion such as chloride or bromide.

The comonomer or comonomers which can be used include: acrylamide, methylacrylamide, diacetone
15 acrylamide, N-alkylated acrylamide and methacrylamide, lower alkyl acrylates or methacrylates, vinylpyrrolidone and vinyl esters; these can be grafted and or crosslinked, such as those described in French Patent
2.189.434.

Further examples which may be mentioned are:

the copolymers of acrylamide and β-methacryloyloxyethyl-trimethylammonium methosulphate which are sold under the names Reten 205, 210, 220 and 240 by the Société Herculès; and the aminoethyl acrylate phosphate/acrylate copolymers which are sold under the name Catrex by National Starch, and also the products described in U.S. Patent 3,372,149 or the polymers referred to as Quaterniums in the Cosmetic Ingredient Dictionary published by the Cosmetic Toiletry and Fragrance Association Inc. Quaterniums in the Cosmetic Ingredient Dictionary.

25 (3) cationic polymers which are:

more hydroxyl and/or carboxyl groups.
c) the quaternary ammonium salts and the products resulting from the oxidation of the polymers of the formulae (III) and (IV) indicated above under a) and b), generally one in which at least one tertiary amino

group in A has been converted into an amine oxide group.

The polymers of the formula (IV) and a process for their preparation are described in French Application 2,280,361.

The polymers of the formula -A-Z-A-Z- (III) can be prepared as indicated in French Patent 2,162,025. 50 (4) the quaternary polyammonium compounds of the formula:

in which R₁, R₂, R₃ and R₄, which are identical or different, represent aliphatic, alicyclic or arylaliphatic radicals containing a maximum of 20 carbon atoms, or lower hydroxyaliphatic radicals, or alternatively R₁ and R₂, and/or R₃ and R₄, together form with the nitrogen atom to which they are attached, a heterocyclic ring optionally containing a second hetero-atom other than nitrogen, or alternatively R₁, R₂, R₃ and R₄ independently represent a group:

60

NSDOCID: <GB 2039938A 1 :

15

20

25

30

35

40

45

50

55

60

65

 R'_3 denoting hydrogen or lower alkyl and R'_4 denoting -CN, or

5

 R'_{5} denoting lower alkyl, R'_{6} denoting hydrogen or lower alkyl, R'_{7} denoting alkylene and D denoting a quaternary ammonium group, A and B represent polymethylene groups which contain from 2 to 20 carbon atoms, can be linear or branched and saturated or unsaturated, and can contain. in the main chain, one or more aromatic rings, to provide a group such as _cH2____or one or more groups Y to provide a group -CH2-Y-CH2-, in which Y denotes O, S, SO, SO2,

$$R'_9$$
 O $||$
25 $\oplus ||$ -S-S-,-N-,-N --X \oplus ,-CH-,-NH-C-NH-,
 $||$ $||$ $||$ R' $_8$ R' $_9$ OH

45

X[©] denoting an anion derived from an inorganic or organic acid, R'₈ denoting hydrogen or lower alkyl and R'₉ denoting lower alkyl, or alternatively A and R₁ and R₃ form a piperazine ring together with the two nitrogen atoms to which they are attached, and B can also denote a group:

 $-(CH_2)_nCO-D-OC-(CH_2)_n$ -, in which D denotes: 40 a) a glycol radical of the formula -O-Z-O, in which Z denotes a linear or branched hydrocarbon radical or a group corresponding to the formula:

$$- \{ cH_2 - cH_2 - 0 \}_{x} - CH_2 - CH_2 - 0r - 0$$

$$- \{ cH_2 - CH_2 - 0 \}_{x} - CH_2 - CH_2 - CH_2 - CH_3$$

$$- \{ cH_3 - CH_2 - 0 \}_{x} - CH_2 - CH_3 - CH_$$

in which x and y denote an integer from 1 to 4, representing a particular degree of polymerisation (in a given molecule), or any number from 1 to 4, representing a mean degree of polymerisation (in the product);

b) a bis-secondary diamine radical, such as a piperazine derivative of the formula:

c) a bis-primary diamine radical of the formula: 50 -NH-Y-NH-, in which Y denotes a linear or branched hydrocarbon radical or the divalent radical -CH2-CH2-S-S-CH2-CH2; or

d) a ureylene group of the formula -NH-CO-NH-, X is an anion, such as chloride or bromide, and n is such that the molecular weight is between 1,000 and 100,000.

Polymers of this type are described, in particular, in French Patents Nos. 2,320,330 and 2,270,846, French 55 Applications Nos. 76/20,261 and 2,336,434 and United States Patents Nos. 2,273,780, 2,375,853, 2,388,614, 2,454,547, 3,206,462, 2,361,002 and 2,271,378, which are hereby incorporated by reference.

Other polymers of this type are described in United States Patents Nos. 3,874,870, 4,001,432, 3,929,990, 3,966,904, 4,005,193, 4,025,617, 4,025,627, 4,025,653, 4,026,945 and 4,027,020, which are hereby incorporated by reference.

60 (5) optionally alkylated, crosslinked polyaminoamides which are water-soluble and can be obtained by cross-linking a polyamino-polyamide (A), prepared by the polycondensation of an acid compound with a polyamine. The acid compound is, for example, (i) an organic dicarboxylic acid, (ii) an aliphatic mono- or di-carboxylic acid having an ethylenic double bond, (iii) an ester of the abovementioned acids, preferably the esters of lower alkanols having from 1 to 6 carbon atoms, and (iv) mixtures of two or more of these

65 compounds. The polyamine may be a bis-primary or mono-or di-secondary polyalkylene-polyamine. Up to

	40 or 50 mol % of this polyamine can be replaced by a bis-primary amine, preferably ethylene-diamine, or by a bis-secondary amine, preferably piperazine, and up to 20 mol % can be replaced by hexamethylene-diamine. The crosslinking can be effected by a crosslinking agent (B) which is an epihalogenohydrin,	
	diepoxide, dianhydride, unsaturated anhydride or bisunsaturated derivative, the crosslinking is generally	
5	carried out using 0.025 to 0.35 molecule of crosslinking agent per amine group of the polyaminopolyamide	5
Ť	(A), preferably from 0.025 to 0.2 and in particular from 0.025 to 0.1 molecule of crosslinking agent per amine	•
•	group of the polyamino-polyamide (A). These polymers and their preparation are described in greater detail	
	in French Patent Application No. 2,252,840.	
	Such crosslinked polymers are perfectly soluble in water at a concentration of 10% by weight without	
10	forming a gel, and the viscosity of a 10% by weight solution in water at 25°C is at least 3 centipoises and	10
ıņ	usually from 3 to 200 centipoises.	
	The crosslinked and optionally alkylated polyaminoamides do not contain any reactive groups, do not	
	have any alkylating properties and are chemically stable.	
	The polyaminoamides (A) themselves can also be used in the compositions of this invention.	
15	(6) the water-soluble, crosslinked polyaminoamides obtained by crosslinking a polyaminoamide, (A,	15
	described above) by means of a crosslinking agent which is:	
	(I) a compound which is (1) a bis-halogenohydrin, (2)a bis-azetidinium compound, (3)a bis-halogenoacyl	
	derivative of a diamine, and (4)a bis-(alkyl halide);	
	(II) an oligomer obtained by reacting a compound (a), which is (1)a bis-halogenohydrin, (2)a bis-	
20	azetidinium compound, (3)a bis-halogenoacyl derivative of a diamine, (4)a bis-(alkyl halide), (5) an	20
	ephihalogenohydrin, (6) a diepoxide, or (7) bis-unsaturated derivative, with a compound (b) which is a	
	difunctional compound which is reactive towards the compound (a); and	
	(III) the product resulting from the quaternisation of a compound (a) or an oligomer (II) and contains one or	
	more tertiary amine groups which can be totally or partially alkylated with an alkylating agent (c), preferably	
25	methyl or ethyl chloride, bromide, iodide, sulphate, mesylate or tosylate, benzyl chloride or bromide,	25
	ethylene oxide, propylene oxide or glycidol, the crosslinking generally being effected with 0.025 to 0.35	
	molecule, in particular 0.025 to 0.2 molecule and more particularly 0.025 to 0.1 molecule, of crosslinking	
	agent per amine group of the polyaminoamide.	
	These crosslinking agents and these polymers, and also the process for their preparation, are described in	
30	French Application No. 2,368,508 which is hereby incorporated by reference.	30
•	(7) the water-soluble polyaminoamide derivatives resulting from the condensation of polyalkylene-	
	polyamines with polycarboxylic acids, followed by alkylation with a difunctional agent, such as adipic	
	acid/dialkylaminohydroxyalkyl-dialkylenetriamine copolymers, in which the alkyl radical contains 1 to 4	
	carbon atoms and preferably denotes methyl, ethyl or propyl, which are described in French Patent	
35	1,583,363.	35
	Compounds which make it possible to obtain valuable results are the adipic acid/dimethylaminohydroxy-	
	propyl-diethylenetriamine copolymers sold under the name Cartaretine F, F_4 or F_8 by SANDOZ.	
·	(8) the polymers obtained by reacting a polyalkylenepolyamine, containing two primary amine groups and	
	at least one secondary amine group, with a dicarboxylic acid which is diglycolic acid or a saturated aliphatic	•
40	dicarboxylic acid having 3 to 8 carbon atoms, the molar ratio of the polyalkylenepolyamine to the	40
	dicarboxylic acid being from 0.8 : 1 to 1.4 : 1, and reacting the resulting polyamide with epichlorohydrin in a	
	molar ratio of epichlorohydrin to the secondary amine group of the polyamide of from 0.5: 1 to 1.8: 1; these	
	polymers are disclosed in U.S. Patents, 3,227,615 and 2,961,347, which are hereby incorporated by reference.	•
	Particularly valuable polymers are those which are sold under the name HERCOSETT 57 by Hercules	
45	Incorporated and have a viscosity of 30 cps at 25°C in a 10% by weight aqueous solution, and those sold	45
	under the name PD 170 or DELSETTE 101 by Hercules, which is an adipic acid/expoxypropyldiethylenet-	
	riamine copolymer.	
	(9) polyalkyleneimines, in particular the polyethyleneimines described in greater detail in U.S. Patent	
	2,182,306, 2,553,696, 2,806,838 and 2,208,085, and also the alkylated or alkoxylated derivatives described in	
50	U.S. Patent 2,039,151 and French Patent 1,506,349.	50
	Amongst the polyethyleneimines and their derivatives, there may be mentioned the products sold under	
	the names PEI 6, PEI 12, PEI 18, PEI 300, PEI 600, PEI 1200, PEI 1800 and PEI 600 E, the last being a	
	polyethyleneimine alkylated with ethylene oxide in a ratio of 1:0.75, and the names TYDEX 14 and TYDEX 16,	
	the latter having a density of about 1.06 and a viscosity at 25°C of more than 1,000 cps. These	er.
- 55	polyethyleneimines are sold by DOW CHEMICAL. The various patents mentioned above are hereby	55
	incorporated by reference.	
	Other polyethyleneimines which can be used according to the invention are those sold by BASF under the	
	name POLYMIN P, which has a density d ₂₀ of about 1.07 and a Brookfield viscosity of 10,000-20,000 in a 50%	
	strength aqueous solution (at 20°C and 20 rpm), POLYMIN SN, which has a density of dd22 ₀ of about 1.06	60
60	and a viscosity of 800-1,800 cps in a 20% strength aqueous solution, and POLYMIN HS, which has a density	60
	d ₂₀ of about 1.07 and a viscosity of 500-1,000 cps in a 20% strength aqueous solution.	
	The products resulting from the reaction of polyethyleneimine with ethyl formate, described in French	
	Patent 2,167,801, can also be used.	

(10) the water-soluble polymers resulting from the condensation of a polyamine and epichlorohydrin, such as the product resulting from the condensation of tetraethylenepentamine and epichlorohydrin.

20

25

35

45

50

55

60

65

10

15

30

(11) the quaternary polyureylenes of the type described in Belgian Patent 77/3,892.

Amongst the non-ionic surface-active agents which are preferably used in the compositions according to the invention, there may be mentioned the products resulting from the condensation of a monoalcohol, an alpha-diol, an alkylphenol or an alkanolamide, such as diglycolamide, with glycidol, such as the products corresponding to the formula: R₄-CHOH-CH₂-O-(CH₂-CHOH-CH₂-O-)_pH in which R₄ denotes an aliphatic, cycloaliphatic or arylaliphatic radical preferably having 7 to 21 carbon atoms, and mixtures thereof, it being possible for the aliphatic chains to contain ether, thioether or hydroxymethylene groups, and in which p is a number from 1 to 10 inclusive, which compounds are described in French Patent 2,091,516; products corresponding to the formula:

R₅O{C₂H₃O(CH₂OH)}₄H,

in which $\rm R_5$ denotes an alkyl, alkenyl or alkylaryl radical and q has a value from 1 to 10 inclusive; and products corresponding to the formula:

 $R_6CONH-CH_2-CH_2-O-CH_2-CH_2-O-CH_2-CHOH-CH_2-O-H_1$

in which R₆ denotes a linear or branched, saturated or unsaturated, aliphatic radical, or a mixture of such radicals, which can optionally contain one or more hydroxyl groups, which have from 8 to 30 carbon atoms and can be of natural or synthetic origin, and r denotes an integer or decimal from 1 to 5. It will be appreciated that p q and r generally represent an average value corresponding to the average degree of condensation since the condensation usually results in the production of compounds with different chain lengths.

Other compounds belonging to this class are polyoxyethyleneated alcohols or alkylphenols or esters of polyethylene glycol or polyglycerol having a linear or branched fatty chain containing 8 to 18 carbon atoms. There may also be mentioned copolymers of ethylene oxide and propylene oxide, products resulting from the condensation of ethylene oxide and propylene oxide with fatty alcohols, polyoxyethyleneated fatty amides, polyoxyethyleneated fatty amines, polyoxyethyleneated fatty acid esters of sorbitol and polyoxyethyleneated fatty acid esters of sucrose.

Amongst these non-ionic surface-active agents, those which are more particularly preferred correspond to the formula: R₄-CHOH-CH₂-O-(CH₂-CHOH-CH₂-O)_pH, in which R₄ denotes a mixture of alkyl radicals having 9 to 12 carbon atoms and p has a statistical value of about 3.5.

 $R_5O\{C_2H_3O(CH_2OH)\}_qH$,

in which $R_{\rm 5}$ denotes $C_{\rm 12}H_{\rm 25}$ and q has a statistical value of 4 to 5, or

R₆-CONH-CH₂-CH₂-O-CH₂-CH₂-O[CH₂CHOH-CH₂O],H,

40 in which R₆ denotes a mixture of radicals derived from lauric, myristic and oleic acids and copra acids and r has a statistical value of 3 to 4.

The preferred oxyethyleneated or polyglycerolated fatty alcohols are polyoxyethyleneated oleyl alcohol containing about 10 mols of ethylene oxide, oxyethyleneated lauryl alcohol containing about 12 mols of ethylene oxide, cetyl alcohol oxyethyleneated with 6 to 10 mols of ethylene oxide, cetyl/stearyl alcohol oxyethyleneated with 3 to 10 mols of ethylene oxide, stearyl alcohol containing 2, 10, 15 or 20 mols of ethylene oxide, oxyethyleneated nonylphenol containing about 9 mols of ethylene oxide, oxyethyleneated octylphenol containing about 5.5 mols of ethylene oxide, polyglycerolated oleyl alcohol containing about 4 mols of glycerol, synthetic C₉-C₁₅ fatty alcohols polyoxyethyleneated with 3 to 12 mols of ethylene oxide, polyoxyethyleneated sorbitan monolaurate containing about 20 mols of ethylene oxide, and the products resulting from the polycondensation of ethylene oxide and propylene glycol.

Amongst the surface-active agents which can be used which contain one or more carboxyl or carboxylate groups in addition to non-ionic groups, there may be mentioned, in particular, polyglyceryl carboxylates, and the carboxylic acids of polyglycol ether corresponding to the formula Alk- $\{OCH_2-CH_2\}_n$ - $\{OCH_2-CO_2H\}$, in which the substitutent Alk corresponds to a linear aliphatic chain having from 12 to 18 carbon atoms and in which n is an integer from 5 to 15, these compounds being in the form of the free acid or of their salts, and in particular the products sold under the name AKYPO RLM 100 by CHEMY, such as the product of the formula $R-\{OCH_2CH_2\}_nOCH_2COOH$, in which R is a mixture of $C_{12}-C_{14}$ alkyl radicals and x is equal to 10.

In particular, good effects can be obtained on dyed or bleached hair, the combination according to the invention in fact making it possible to recover the appearance and the condition of natural hair.

The compositions according to the invention are preferably aqueous and can be used as such for the purpose of treating the various fibrous materials mentioned above. However, they can also contain organic solvents such as alkanols having from 1 to 8 carbon atoms, such as ethanol, isopropanol, benzyl alcohol, phenylethyl alcohol and methoxy-, ethoxy-, propoxy- and butoxyethanol, alkylene glycols, such as ethylene glycol, propylene glycol, butylene glycol and diethylene glycol monoethyl ether, and also esters, such as the

2.14%

0.96%

12.5%

4%

acetate of ethylene glycol monomethyl ether or monoethyl ether and the esters of fatty acids and lower alcohols, such as isopropyl myristate or palmitate. These solvents are generally present in an amount from 0.5 to 30% by weight. The compositions can contain, in addition to the abovementioned ingredients, adjuvants which are 5 normally used in compositions for treating textiles or hair. They can contain, in particular, perfumes, dyestuffs, the purpose of which can be to colour either the composition itself or the hair or the textiles, preservatives, sequestering agents, thickeners, emulsifiers, softeners, synergistic agents and foam stabilisers, depending on the application envisaged. The dyestuffs used for dyeing the materials treated include oxidative dyestuffs, such as those of the 10 well-known diamine, aminophenol or phenol types, and direct dyestuffs, such as azo dyestuffs, anthraquinone dyestuffs, nitrobenzene dyestuffs, indamines, indoanilines, indophenols as well as other

oxidative dyestuffs, such as leuco derivatives of these compounds, these various types of dyestuff being used singly or as a mixture.

The compositions according to the invention are preferably used for washing.

For cosmetic use, they can be in the form of, for example, a shampoo, but also in the form of colouring products, rinsing lotions to be applied before or after shampooing, before or after colouring or bleaching or before or after permanent waving, brushing lotions, styling and restructuring lotions and gels.

The perfumes which can be used in these compositions are cosmetically acceptable perfumes; they are generally present in an amount from 0.1 to 0.5% by weight.

Alkalising or acidifying agents such as acetic, lactic, citric and phosphoric acids and ammonia and mono-di-or tri-ethanolamines can be added to the compositions.

The following Examples further illustrate the present invention; the percentages are indicated by weight unless otherwise stated.

The anionic polymers are 100% neutralised with sodium hydroxide.

25

The following composition is prepared:

Anionic polymer sold under the name

30 GANTREZ ES 425

Cationic polymer referred to as PAA-R2

Non-ionic surface-active agent referred

35 to as TA-1

NaCi

40

50

Water q.s.p.

The ratio of equivalents of cationic units to equivalents of anionic units is 3 and the pH is equal to 6.

This homogeneous composition is used as a shampoo; when applied to dirty hair after wetting, it produces a gentle foam.

When wet, the hair is easy to comb out and soft to the touch.

When dry, the hair possesses a good hold, has a smooth and soft feel and is very springy, very bulky and 45 very manageable.

After styling, the hair style is firm and holds well and the hair is glossy.

Similar results are obtained with the compositions illustrated in Table I which follows:

TABLE I

50

25

30

35

40

55	Exam- ple	Anionic polymer	%	Cationic polymer	%	Surface- active agent	%	Alkali metal salt	%	Cationic polymer/ anionic polymer equivalents	55
	2	GANTREZ ES 425	2.04	PAA - R2	0.96	TA-1	12.5	NaCl	4	0.5	
60	3	GANTREZ ES 425	1.56	PAA - R2	1.44	TA-1	12.5	NaCl	4	· 1	60
	.4	GANTREZ ES 425	1.57	PAA - R2	1.43	TA-1	12.5	NaCl	6	1	

15

20

1.97%

1.03%

12.5%

2%

A shampoo having the following composition is prepared:

Anionic polymer sold under the name

Cationic polymer referred to as PAA-I

Non-ionic surface-active agent referred

Water gsp 100

The pH is equal to 6 and the ratio of cationic polymer/anionic polymer in equivalents is equal to 0.66. When applied to dirty hair after wetting, the formation of a gentle foam is observed.

When wet, the hair is easy to comb out.

When dry, the hair possesses a good hold and is springy and the head of hair is bulky.

The hair style holds well and the hair is soft.

Similar results are obtained on modifying the proportions of the various polymers in the following

TABLE !!

25 Alkali % Cationic % Surface Cationic % Exam- Anionic polymer polymer polymer/ active metal ple anionic agent salt polymer 30 30 equivalents 12.5 NaCl 4 1 1.33 TA-1 1.67 PAA-I 6 **GANTREZ ES 425** 3 35 4 0.89 PAA-I 2.1 TA-1 12.5 NaCi **GANTREZ ES 425** 35 7 NaCl 0.5 0.85 TA-1 12.5 **GANTREZ ES 425** 2.15 PAA-I 8

The pH of the compositions of Examples 6 and 7 is equal to 5.9 and 5.7 respectively. 40

Examples 9 to 62

Table III which follows is intended to illustrate other methods of carrying out the present invention.

This table indicates the nature of the anionic polymer, the cationic polymer, the surface-active agent and 45 the alkali metal salt, and the ratio of equivalents of cationic units to equivalents of anionic units. As in the preceding tables, the water added in all cases to make up 100 g is not included in this table.

The compositions used as a shampoo gave similar results to those reported above, in particular as regards the ease of comb-out and the softness of the hair when wet, and the springiness and softness of the hair when dry, and the hair style possesses bulk, firmness and a good hold. All these compositions are 50 homogeneous and stable under normal storage conditions.

50

TABLE III

Exam- PLE	PC	OLYMER			Surface- Active		Alkali Metal		CATIONIC Polymer/
No.	Anionic .	%	Cationic	%	Agent	%	Salt	%	Anionic polymer Equiva- lents
9	GANTREZ ES 425	2.5	AZA - 1	0.5	TA - 1/8	1/81/4.5/8	NaCl	4	0.5
10	GANTREZ ES 425	2.16	AZA - 1	0.84	TA-1	12.5	NaCl	4	•1
11	GANTREZ ES 425	1.9	PAA - R1	1.1	TA - 1	12.5	NaCl	4	0.5
12	GANTREZ ES 425	1.4	PAA - R1	1.6	TA - 1	12.5	NaCl	4	.1
13	GANTREZ ES 425	0.68	PAA - R1	2.32	TA - 1	12.5	NaCl	4	3
14	GANTREZ ES 425	0.89	PAQ-1	2.1	TA - 1	12.5	NaCl	4 -	3
15	GANTREZ ES 425	2.15	PAQ-1	0.85	TA - 1	12.5	NaCl	4	0.5
16	GANTREZ ES 425	1.68	PAQ-1	1.32	TA - 1	12.5	NaCl	4	1
17	GANTREZ ES 425	1.69	PAA - RA -1	1.3	TA - 1	12.5	NaCl	4	0.5
18	GANTREZ ES 425	1.18	PAA - RA -1	1.82	TA - 1	12.5	NaCl	4	1
19	GANTREZ ES 425	0.53	PAA - RA -1	2.47	TA - 1	12.5	NaCl	4	3
20	GANTREZ ES 425	1.57	PAA - RA 2	1.43	TA - 1	12.5	NaCl	4	0.5
21	GANTREZ ES 425	1.06	PAA - RA 2	1.94	TA - 1	12.5	NaCl	4	1
22	GANTREZ ES 425	0.46	PAA - RA 2	2.54	TA - 1	12.5	NaCl	4	3 .
23	GANTREZ ES 425	1.5	PAQ-2	1.5	TA - 1	12.5	NaCi	4	3
24	GANTREZ ES 425	2.12	PAQ-2	0.88	TA - 1	12.5	NaCl	4	1
25	GANTREZ ES 425	1.88	AZA - 2	1.12	TA-1	12.5	NaCl	4	0.5
26	GANTREZ ES 425	1.36	AZA - 2	1.84	TA - 1	12.5	NaCl	4	1
27	GANTREZ ES 425	0.66	AZA - 2	2.34	TA - 1	12.5	NaCl	4	3
28	GANTREZ ES 425	0.46	PAA - RA 3	2.54	TA - 1	12.5	NaCl	4	3
29	GANTREZ ES 425	1.06	PAA-RA3	1.94	TA - 1	12.5	NaCl	4	1
30	GANTREZ ES 425	1.57	PAA-RA3	1:43	TA - 1	12.5	NaCl	4	0.5
31	GANTREZ ES 425	0.46	PAA - RA 3	2.54	TA - 1	12.5	NaCl	2	3 .
32	GANTREZ ES 425	1.71	PAA-2	1.29	TA - 1	12.5	NaCl	4	0.5
33	GANTREZ ES 425	1.19	PAA-2	1.81	TA - 1	12.5	NaCl	4	1
34	GANTREZ ES 425	0.54	PAA-2	2.46	TA - 1	12.5	NaCl	4	. 3
35	SMA 1,000	1.39	PAQ - 1	1.61	TA - 2	10	NaCl	4	1
36	SMA 1,000	0.56	PAQ - 1	2.42	TA - 2	10	NaCl	4	3

Exam- ole		POLYMER			Surface Active		Alkali metal		Cationic Polymer
No.	Anionic	%	Cationic	%	Agent	%	Salt	%	Anionic polymer
	•		•					•	Equiva- lents
7	SMA 1,000	1.10	AZA - 1	1.9	TA - 2	10	NaCl	4	0.4
8	SMA 1,000	1.76	AZA - 1	1.24	TA - 2	10	NaCl	4	1
9	SMA 1,000	1.76	AZA-1	1.24	Polyethylene glycol lauryl ether (11-120E)	10	NaCl	4	1
	SMA 1,000	1.1	PAA-1	1.9	polyethylene glycol lauryl ether (11-120E)	10	NaCl	4	1
11	SMA - 1,000	0.64	PAA - RA-2	2.36	"	10	NaCl	4 .	1
2	SMA - 1,000	0.68	PAA - RA-3	2.32		10	NaCl	4	. 1
3	SMA - 1,000	1.7	PAQ - 2	1.3	"	10	NaCl	4 ,	1
4	Aristoflex A	2.7	AZA - 1	0.3	Surfactant 10 G	. 10	NaCl	4	1
5	Aristoflex A	1.96	PAA - RA-3	1.04	"	10	NaCl	4	1
6	Aristoflex A	2.38	PAA - ra-3	0.62	"	10	NaCl	4	0.5
7	Aristoflex A	1.96	PAA - RA-3	1.04	"	10	NaCl	4	1
8 ,	SMA - 1,000	1.2	PAQ - 1	1.76	"	10	NaCl	4	1
9	GANTREZ ES 425	1.54	PAA - 1	1.46	TA - 3	10	NaCl	4	1
0	VERSICOL K 11	1.42	PAQ-2	1.56	TA - 1	10	NaCl	4	1
1	28 - 29 - 30	2.38	PAA-1	0.62	Surfactant 10 G	10	NaCl	4	1
2	Aristoflex A	1.92	PAA - RA 2	1.08		10	NaCl	. 4	1
3	Aristoflex A	2.44	PAQ-1	0.56		10	NaCl	4	1
4	Aristoflex A	2.68	PAQ - 2	0.32	"	10	NaCl	4	1
5	GANTREZ ES 425	1.06	PAA-RA3	1.94	TA - 5	10	NaCl	4	1
6	GANTREZ ES 425	1.5	PAA-1	1.5	BRIJ 35	10	NaCl	4	1
7	GANTREZ ES 425	1.05	PAA-RA3	1.95	BRIJ 35	10	NaCl	4	1
8	ARISTOFLEX A	2.36	PAA - 1	0.64	Surfactant 10 G	10	NaCl	4	1

					,		-	•	. '	
5	9	GANTREZ ES 425	2.12	PAQ-2	0.88	TA-3	10	NaCl	4	1
5 6	iO	VERSICOL K 11	1.44	PAQ-2	1.56	TA - 1	10	KCI	4,	1 5
6	i1	28 - 29 - 30	2	PAA-RA3	1	Surfactant 10 G	. 10	LiCl	4	1
10 6	52	HYDAGEN F	2.5	Polyquart H	0.5	AKYPO RLM 100	10	NaCl	4	0.2 10
		ple 63 following composition	n is prepa	red:						15
		ic polymer sold under ICOL K 11	the name				1%			15
	Cation	nic polymer referred to	as PAQ -	3			2%			20
20	Surfac	ce-active agent referred	d to as TA	-1			10%			20
ı	NaCl					•	4%			
25 V	Vater	qsp 100	•		-	•				25
30 s	wet ar Who This Sim surfac	s composition is used and the hair is easy to co en dry, the hair has a go s composition can also illar results are observe e-active agent referred wise being the same.	omb out. ood hold be used f ed on repl	and a soft feel a or restructuring acing the surfa	and is sp g the ha ce-activ	oringy and very m ir. e agent referred (nanageat to as TA ·	ole. - 1 by the		30
· i	Tab nven	ples 64 to 71 le IV which follows is in tion; these composition same way as above.	ntended to ns are ho	o illustrate othe mogeneous an	er comp d impar	ositions falling w t advantageous c	rithin the cosmetic	scope of t properties	he s to the ha	35 ir

TABLE IV

	Exam- ole No.	POLYMER Anionic	%	Cationic	%	Surface- Active Agent	%	Alkali Metal Salt	Cationio	Polymer/ Anionic Polymer Equivalents	
)	64	HYDAGEN F	1.8	PAQ-3	1.2	TWEEN 20	10	NaCl	4	0.44	. 1
	65	HYDAGEN F	2	PD 170	1	TA-1	8	NaCl	4	0.44	
5				·		GLUCAMAT SSE 20	2				1
	66	HYDAGEN F	1.7	GAFQUAT 755	1.3	TWEEN 20	10	NaCl	4	0.11	
0	67	VERSICOL K 11	2	PAQ-3	1 .	TA-1	10	NaCl	4	0.24	2
٠.	68	VERSICOL K 11	2	PAQ-3	1.	AKYPO RLM 100	10	NaCl	4	0.25	
5	69	VERSICOL E 5	0.5	CATREX	2.5	AKYPO RLM	10	NaCl	4		:
•	Examp	ole 70	ing 127	o/m² and cons	sistina o	100	/cm and	17 weft	picks/cm	, is scoured	
0	A wo with m A sai	oollen cloth, weigh ethylene chloride. mple of this fabric				f 20 warp yarns		osition:	-	, is scoured	
	A wo with m A sai	oollen cloth, weigh ethylene chloride. mple of this fabric eset 57	is imme	rsed in a solut		f 20 warp yarns			-	, is scoured	
	A wo with m A sai Hercos	oollen cloth, weigh ethylene chloride. mple of this fabric	is imme	rsed in a solut		f 20 warp yarns		osition:	-	, is scoured	
5	A wo with m A sai Hercos Versico 100% v	oollen cloth, weigh ethylene chloride. mple of this fabric sset 57 ol E5 (neutralised t	is imme	ersed in a solut tent of containing 30 m	tion havi	f 20 warp yarns		osition: 0.5	g	, is scoured	
5	A wo with m A sai Hercos Versico 100% v	pollen cloth, weigh tethylene chloride. Imple of this fabric isset 57 Tol E5 (neutralised t with NaOH)	is imme	ersed in a solut tent of containing 30 m	tion havi	f 20 warp yarns		osition: 0.5 1 g	g	, is scoured	
5	A wo with m A said Hercos Versica 100% v Oxyeth of ethy NaCl Water	pollen cloth, weigh tethylene chloride. Imple of this fabric sset 57 of E5 (neutralised t with NaOH) hyleneated nonylp ylene oxide (per mo	is imme o the ex henol co	ersed in a solute tent of containing 30 menol)	tion havi	f 20 warp yarns	g comp	0.5 1 g 1 g 4 g 100 g	g		
5	A wo with m A said Hercos Versica 100% v Oxyeth of ethy NaCl Water	pollen cloth, weigh ethylene chloride. mple of this fabric set 57 ol E5 (neutralised t with NaOH) nyleneated nonylp ylene oxide (per mo	o the ex henol cool of phe	ersed in a solut stent of containing 30 m enol)	nols o of equi	f 20 warp yarns ng the followin	g comp	1 g 1 g 4 g 100 g	g valents c	of anionic units	•

5	ambient of water 45° stren 2 ml o	e 71 ple of woollen cloth temperature using and 0.1 ml of analy tigth of active chlori f sodium bisulphite . This sample is imr	a lique tical gr ne are soluti	or ratio of 1/1 rade HCl. Aft added and tl on are then a	00.1g er an ir his is al added a	of scoured mpregnatio llowed to ac and rinsing	wool is i n time of at for 30 i with run	ntroduced i f 5 minutes, minutes. ning water	into a mi , 4 ml of	xture of 100 ml Javelle water of	5
	Polymer	referred to as PAQ	-3					1	g		10
10	Hydageı	n F					J	. 1	g		
	Surface-	active agent TA-1				•		1	g		
15	NaCl							5	5 g		15
	Water q	sp				·		100) g		
20	The ra	H is adjusted to 7 water to of equivalents of treatment, the samp	f catio	nic units to e	equival by a v	ents of anio ery marked	nic units stiffness	s is equal to s.	0.7.		20
	Example water.	e 72 Ollen fabric, weighir e 71, is neutralised v ample is immersed	with a	20% strength	n soluti	on of sodiu	m carbo	nate and th	hlorinat en rinse	ed as indicated in d with running	25
	MERQU	·	•			-			1 g		
30		OL E5 (neutralised t y NaOH)	o the e	extent of					1.5 g	•	30
35		yleneated nonylphe ene oxide	nol co	ntaining 30 r	nols				1 g		35
	NaCl	. •							4 g		
40	Water q	sp oH is adjusted to 7 w	ith HC	·				10	0 g		40
45	The range A gre Simil compose	atio of equivalents of ater stiffness of the ar results to those notions using a liquo of about 70% and of also be effectively.	of cation wool to the month of	nic units to o reated is also ned above an of 1/40 in a b ally dried in a	o obse re obta oath wl in over	rved in this ined on imp hich can be n at 60°.	case. oregnati hot or co	ng the fabri old. The fab	ic in the oric is sq	various ueezed off to a	45
50	On w	les 73 to 76 rashing the treated v sitions;	vool, a	s mentioned	l in Exa	amples 70 to	72, by	means of th	ne follow	ring	50
55	Exam- ple	Anionic Polymer	%	Cationic Polymer	%	Surface- active agent	%	Alkali metal salt	%	CP/AP Equivalents	55
	73	Gantrez ES 425	2.5	AZA - 1	0.5	TA-1	12.5	NaCl	. 4	0.5	
60	74	SMA 1,000	1.10	AZA - 1.	1.9	TA - 2	10	NaCl	4	0.4	60
	75	GANTREZ ES 425	0.46	PAA - RA3	2.54	TA - 1	12.5	NaCl	4	3	
6 F	76	VERSICOL K 11	1.44	PAQ-2	1.56	TA - 1	10	KCI	4	1 .	65
65											

	and on rinsing with running water, an increase in	n the firmness of the wool is	observed.	÷
5	Examples 77 to 82 These Examples illustrate the treatment of fibraqueous composition containing (1) a cationic pa surface-active agent.	res and fabrics, as indicated l olymer, (2) an anionic polym	below, with a homogenous ner, (3) an alkali metal salt and (4)	5
	Fibres:	· .		
10	Polypropylene 2.5 D Montedison Polyester CHEROTAN 3 D/60			10
	Fabrics:		•	
15	White test fabric	NYLON	(Polyamide 6.6)	15
	White test fabric	TERGAL	(Polyester)	
	White test fabric	CRYLOR	(Polyacrylonitrile)	20
20	White test fabric	THERMOVYL	(Polyvinylchloride)	
	White test fabric	RILSAN	Polyamide 11)	
25	White test fabric	TRIACETATE		25
	White test fabric	VISCOSE	•	
30	Fabric	100% COTTON	(weight 163g/m², 29 wefts per cm, 52 yarns per cm).	30
	Fabric	55% wool	(weight 324g/m²,	
35		45% poly- ester	20 wefts per cm, 35 yarns per cm).	35
	Before the treatment, the fibres and fabrics w (Coptal BR sold by UGINE KUHLMAN), rinsed of temperature.	ere washed with a solution o opiously with running water	containing 2% of detergent then dried at ambient	40
40 4!	Application of the compositions: The test fibres and fabrics were immersed in a for 15 minutes. Part of the test materials were risqueezed out between a series of rollers and the ln some cases the drying was followed by a the composition of the cases the drying was followed by a the case was a followed by a followed by a the case was a followed by a fol	nsed and the other part was en the test materials were dr hermal treatment at a tempe	not rinsed but excess liquid was ried.	45
	compatible with the properties of the materials	and polymers used.		
	Example 77			
5	PAA-RI		3 g	50
	GANTREZ ES 425 completely neutralised with NaOH		3 g	:
5	5 TAI		1 g	55
	Na Cl		3 g	•
	Water q.s.q.		100 g	60
6	D PH	= 8	•	
	•			

Fabrics

The fabrics were immersed in the composition for 15 minutes at ambient temperature with a bath ratio of 40:1. Half the test materials were then rinsed with running water and the other half were not rinsed but

squeezed out (degree of squeezing out 70%). All the fabrics were dried in an apparatus with forced air at 60°C for 30 minutes.

The following results were observed:

		٠.	
5	100% Cotton fabric:		5
	With or without rinsing an increase in the rigidity of the fabric was noted. The effect was more marked		
	however without rinsing. This rigidity is superior to that which is achieved using each of the polymers		
	separately from a solution containing the same products as in the mixture used.		
10	Wool/polyester fabric:	1	0
	Same results as above although the increase in the rigidity was more marked than with the cotton fabric.		
	4000/ Nulsan fahain		
	100% Nylon fabric:		
	Results identical to those above. The effect is the same degree as with cotton.		

15
100% Viscose fabric:
Very clear results and of the same type as those above. The effect is very marked when no rinsing takes place.

20 Crylor fabric: The effect is very marked, more marked than for the nylon and viscose. It is practically the same whether or not the fabric has been rinsed.

Fibres:

25 Cherotan polyester:

Very slight increase in rigidity of a sample of fibres which appear denser and more compact when the

Very slight increase in rigidity of a sample of fibres which appear denser and more compact when the composition is not rinsed.

Polypropylene:
30 Identical results to those obtained on the polyester.
30 30

Example 78

 CARTARETINE F4
 0.5 g

 MERQUAT 100
 0.25g

 VERSICOL E 5
 0.5 g

 Na Cl
 3g
 40

40 Na Cl 3g 40
TA 2 10 g

Water q.s.p. 100 g 45

pH = 9.1 (Na OH)

This composition applied to a 100% Crylor fabric gave a very slight increase in rigidity. The effect is comparable whether or not the material is subsequently rinsed.

5 g

55 MERQUAT 100 2.5 g 55

5 MERCOAT TOU 2.5 g

VERSICOL E 5 0.5 g

Na Cl 5 g

60 60

NIC 33 5 g

Water q.s.p. 100 g

pH = 9.1 (Na OH)

CARTARETINE F4

This composition applied to a 100% Crylor fabric led to a very slight increase in rigidity. The effect is a little more marked when the fabric is not subsequently rinsed.

	Indie Harked When the labite is not outsequently thised.	•		
	Example 80			5
5	PD 170	0.65 g		
	GANTREZ ES 425, 100% neutralised with NaOH	1 g		
10	Na Cl	5 g	•	10
	SANDOPAN DTC acid	5 g		
	Water q.s.p.	100 g		15
15	pH = 8.9 (Na OH)		·	19
20	This composition applied to a 100% cotton fabric led to a very slight increas application was not followed by rinsing. The same result was obtained on dry oven.	e in rigidity when ing the fabric to a	the 100°C in an	20
	Example 81	. :		•
	POLYMIN HS	1 g		25
25	VERSICOL E 5	3 g		
	Na Cl	5 g	• •	
30	AKYPO RLM 100	2.5 g		30
	Water q.s.p.	100 g		
35 40	This composition was applied to wool polyester fabric (55/45) and it gave a when the application was not followed by rinsing. The same result was obtain an oven or subjecting the already dried material to a subsequent thermal trainingtes.	ned on drying the	tabric at 100°C	35 40
	POLYMIN HS	3 g		,
AE	DARVAN No. 7	3 g		45
40	Na Cl	3 g		
	NI 170	1 g		
.50		100 g		50
	pH = 9.5 (Na OH)		*	
55	This composition applied to a 100% tergal fabric gave a very slight increase marked when the application was not followed by rinsing. The various abbreviations and tradenames, used in the above Examples at below.		•	55
	Anionic polymers Gantrez ES 425 Poly-(methyl vinyl ether/maleic acid) monobutyl ester sold by 28 - 29 - 30 Vinyl acetate/crotonic acid/vinyl neodecanoate terpolymer sold by Aristoflex A Terpolymer of vinyl acetate/crotonic acid and polyethylene glycoversicol K 11 Methacrylic acid polymer having a molecular weight of 10,000 525% strength solution, sold by ALLIED COLLOIDS.	y NATIONAL STA ol sold by HOECH	RCH. ST.	60 65

Versical E 5 Mixture of acrylic acid homopolymer and copolymer, having a viscosity of 16 cps in a 25% strength solution and a molecular weight of about 3,500, sold by ALLIED COLLOIDS.

SMA - 1,000 Styrene/maleic anhydride copolymer having a mean molecular weight of 1,600 and a viscosity of 17 cps in 15% strength aqueous ammonia, sold by ARCO CHEMICAL CO.

5 Darvan No. 7 Sodium polymethacrylate sold by VAN DER BILT.

5

10

15

Cationic polymers

AZA - 1: Cationic product resulting from the polycondensation of piperazine, diglycolamine and epichlorohydrin in the molar proportions of 4/1/5, described in Example 2 of French Patent 2,280,301.

0 AZA - 2: Cationic product resulting from the polycondensation of bis-(chloroacetyl)-piperazine and piperazine in equimolecular amounts.

PAA - 1: Product resulting from the polycondensation of adipic acid and diethylenetriamine in equi-molecular amounts.

PAA - 2: Product resulting from the polycondensation, with diethylenetriamine, of the product resulting from the reaction of 2 mols of methyl itaconate and 1 mol of etylenediamine.

PAA - R1: Polymer resulting from the crosslinking of the polymer PAA - 1 with epichlorohydrin (11 molecules of epichlorohydrin per 100 amine groups).

PAA - R2: Polymer obtained by crosslinking the polymer PAA - 1 with a random oligomeric crosslinking agent of the formula

20 $c_1 c_{H_2} - c_{HOH} - c_{H_2} - \frac{1}{2} c_1$ 20

PAA - RA1: Polymer resulting from the alkylation of the polymer PAA - R1 with tert.-butyl glycidyl ether.

PAA - RA2: Polymer resulting from the alkylation, with glycidol, of the polymer referred to as PAA - R1.

25 PAA - RA3: Polymer resulting from the alkylation of the polymer PAA R1 with glycidyltrimethylammonium chloride.

This polymer is prepared in accordance with the following process:

158 g (that is to say 1,000 meq of epoxide) of glycidyltrimethylammonium chloride are added to 1,314 g of PAA - R1 in a 20% strength aqueous solution (1,123 meq of basicity). The mixture is heated for 2 hours at

30 60°C and then diluted with 2 litres of water.

30

The mixture is kept at 60°C for a further 2 hours.

A yellow solution containing 11.8% of active ingredient is obtained.

Characteristics of the solution:

Chlorine number = 0.34 meq/g

35 Base number: 0.29 meq/g

35

Viscosity (in a solution containing 10% of active ingredient) at 25°C: 0.24 poise for a velocity gradient of 14.7 second⁻¹.

PAQ - 1: Polymer having recurring units of the formula

40

$$\begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{2} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
ch_{3} & ch_{3} \\
 & ch_{3}
\end{bmatrix} = \begin{bmatrix}
c$$

45 PAQ-2: Polymer having recurring units of the formula

45

50

55

40

50

PAQ - 3: Polymer having recurring units of the formula

$$= \begin{bmatrix} c_{H_3} & c_{H_3} & c_{H_3} \\ \vdots & c_{H_2} & c_{H_3} & c_{H_3} \\ \vdots & \vdots & \vdots \\ c_{H_3} & c_{H_3} & c_{H_3} \end{bmatrix}_{2 = 0}^{C_{H_3}}$$

55 n being equal to about 6. Gafquat 755: Quaternary vinylpyrrolidone copolymer having a molecular weight of 1,000,000, marketed by GENERAL ANILINE.

MERQUAT 550 : Dimethyldiallylammonium chloride/acrylamide copolymer having a molecular weight > 500,000, sold by MERCK.

60 PD 170: Adipic acid/epoxypropyl-diethylenetriamine copolymer sold by HERCULES, of the formula

2. A composition according to claim 1 which has a pH of 5 to 8.

25

30

40

45

55

60

65

35

40

50

- 3. A composition according to claim 1 or 2, in which the cationic polymer and the anionic polymer are each present in an amount from 0.25 to 3% by weight.
- 4. A composition according to any one of claims 1 to 3 in which the alkali metal salt is present in an amount from 0.25 to 8% by weight.
- 5. A composition according to any one of claims 1 to 4, in which the surface-active agent is present in an amount from 1 to 50% by weight.
- 6. A composition according to any one of claims 1 to 5, in which the ratio of anionic polymer to cationic polymer is from 5:1 to 0.04:1, expressed as the ratio of equivalents of cationic units to the equivalents of anionic units.
- 7. A composition according to any one of claims 1 to 6, in which the alkali metal salt is a potassium, sodium or lithium salt.
 - 8. A composition according to any one of claims 1 to 7, in which the alkali metal salt is a halide, sulphate, acetate or lactate.
- 9. A composition according to any one of claims 1 to 8, in which the anionic polymer contains units derived from an unsaturated mono- or di-carboxylic acid of the formula

$$R_1$$
 $C = C$
 R_3
 $C = C$
 R_3

in which n is 0 or an integer from 1 to 10; A denotes a methylene group which is joined to the carbon atom of the unsaturated group and/or to the adjacent methylene group if n is greater than 1, either directly or via a hetero-atoms; R₁ denotes a hydrogen atom or a phenyl or benzyl group; R₂ denotes a hydrogen atom or an alkyl group of 1 to 4 carbon atoms or a carboxyl group; and R₃ denotes a hydrogen atom, an alkyl group of 1 to 4 carbon atoms, a CH₂ -COOH group or a phenyl or benzyl group.

- 10. A composition according to any one of claims 1 to 9 in which the cationic polymer is a polyamine or
 25 quaternary polyammonium polymer, in which the amine or ammonium group forms part of the polymer chain or is joined to the latter.
 - 11. A composition according to claim 10, in which the cationic polymer is an optionally quaternised vinylpyrrolidone/aminoalcohol acrylate copolymer or a cellulose ether derivative containing quaternary ammonium groups.
 - 12. A composition according to claim 10, in which the cationic polymer is chosen from amongst:
 (1) a water-soluble cyclic polymer having a molecular weight of 20,000 to 3,000,000 and containing chain units of the formula (II) or (II'):

$$(II) = \begin{bmatrix} CH_2 - R^*C & CH_2 \\ H_2C & CH_2 \\ R & + R \end{bmatrix}$$

$$(II') = \begin{bmatrix} CH_2 - R^*C & CH_2 \\ H_2C & CH_2 \end{bmatrix}$$

$$(II) = \begin{bmatrix} CH_2 - R^*C & CH_2 \\ CH_2 & CH_2 \end{bmatrix}$$

$$(II) = \begin{bmatrix} CH_2 - R^*C & CH_2 \\ CH_2 & CH_2 \end{bmatrix}$$

$$(II) = \begin{bmatrix} CH_2 - R^*C & CH_2 \\ CH_2 & CH_2 \end{bmatrix}$$

$$(II) = \begin{bmatrix} CH_2 - R^*C & CH_2 \\ CH_2 & CH_2 \end{bmatrix}$$

in which R" denotes hydrogen or methyl, R and R' independently denote an alkyl group having from 1 to 22 carbon atoms, a hydroxyalkyl group or a lower amidoalkyl group or R and R' together denote, with the nitrogen atom to which they are attached, a heterocyclic group and Y⁻ denotes a bromide, chloride acetate, borate, citrate, tartrate, bisulphate, bisulphate, sulphate or phosphate anion.

45 (2) an acrylic acid or methacrylic acid homopolymer or copolymer comprising the chain unit:

- 55 in which R₁ is H or CH₃; A is a linear or branched alkyl group having 1 to 6 carbon atoms or a hydroxyalkyl group having 1 to 4 carbon atoms; R₂, R₃ and R₄ independently denote an alkyl group having 1 to 18 carbon atoms or a benzyl group; R₅ and R₆ independently denote H or alkyl having 1 to 6 carbon atoms, and X denotes a methosulphate or halide anion.
- (3) a cationic polymer having recurring units of the formula: -A-Z-A-Z- (III), in which A denotes a radical containing 2 amino groups, and Z denotes the symbol B or B' and B and B', which are identical or different, denote a linear or branched alkylene radical which is unsubstituted or substituted by one or more hydroxyl groups and which can contain 1 to 3 aromatic and/or heterocyclic rings as well as oxygen, nitrogen and sulphur atoms; or having recurring units of the formula:
- -A-Z₁-A-Z₁- (IV), in which A is as defined abve and each Z₁ independently denotes B₁ or B'₁ such that at least one Z denotes B'₁, B₁ denotes a linear or branched alkylene or hydroxyalkylene radical having up to 7 carbon

atoms in the main chain and B'1 denotes a linear or branched alkylene radical having up to 7 carbon atoms in the main chain and is unsubstituted or substituted by one or more hydroxyl radicals and contains one or more chain nitrogen atoms, which are substituted by an alkyl chain which optionally contains one or more chain oxygen atoms and one or more hydroxyl and/or carboxyl groups; or a quaternary ammonium salt of a 5 polymer of formula (III) or (IV) or an oxidation product of a polymer of formula (III) or (IV) such that at least one tertiary amino oxide group in A has been converted into an amine oxide group. (4) a quaternary polyammonium compound of the formula:

10

10

in which R_1 , R_2 , R_3 and R_4 , which are identical or different, represent an aliphatic, alicyclic or arylaliphatic radical containing a maximum of 20 carbon atoms or a lower hydroxyaliphatic radical, or R₁ and R₂, and/or 15 R₃ and R₄, together form, with the nitrogen atom to which they are attached, a heterocyclic ring optionally containing a second hetero-atom other than nitrogen, or R₁, R₂, R₃ and R₄ independently represent a group of formula:

15

-CH₂-CH(

20

25 in which R'3 denotes hydrogen or lower alkyl and R'4 denotes -CN,

25

30

35 In which R'_5 denotes lower alkyl R'_6 denotes hydrogen or lower akyl, R'_7 denotes alkylene and D denotes a quaternary ammonium group, A and B independently represent a linear or branched, saturated or unsaturated aliphatic group of 2 to 20 carbon atoms, which contain in the chain one or more aromatic rings, and/or one or more groups of formula - CH₂ -Y-CH₂-, in which Y denotes O, S, SO, SO₂,

35

40 || -- X[©], -CH - , - NH -C- NH - , OH

Δſ

45 or 50

45

R'R

50

In which X^{Θ} denotes an anion of an inorganic or organic acid, R'₈ denotes hydrogen or lower alkyl and R'₉ denotes lower alkyl, or A,R₁ and R₃ form a piperazine ring together with the two nitrogen atoms to which they are attached, and B can also denote a group of formula: - (CH₂)_nCO-D-OC-(CH₂)_n-, inwhich D denotes: a) a glycol radical of the formula -O-Z-O-, in which Z denotes a linear or branched hydrocarbon radical or a group of the formula:

55

60

 $- \left[\text{CH}_{2} - \text{CH}_{2} - \text{O} \right]_{X} - \text{CH}_{2} - \text{CH}_{2} - \text{Or} - 0 \left[\begin{array}{cccc} \text{CH}_{2} - \text{CH} - 0 \\ & & & \\ & & & \\ \text{CH}_{3} \end{array} \right]_{V} - \text{CH}_{2} - \text{CH}_{3}$

60 in which x an y independently denote an integer from 1 to 4.

b) a bis-secondary diamino radical.

c) a bis-primary diamino radical of the formula:

-NH-Y-NH-, in which Y denotes a linear or branched hydrocarbon radical or the divalent radical

-CH2-CH2-S-S-CH2-CH2-; or 65 d) a ureylene group of the formula -NH-CO-NH-; and n is such that the molecular weight is from 1,000 to

15

20

25

30

35

40

45

50

55

0			

(5) a polyaminoamide.

(6) a crosslinked polyaminoamide which is:

a) water-soluble, optionally alkylated, crosslinked polyaminoamide obtained by crosslinking a polyami5 noamide derived from the polycondensation of an acid compound with a polyamine, with a crosslinking
agent which is an epihalogenohydrin, diepoxide, dianhydride, unsaturated anhydride or a bis-unsaturated
derivative, in an amount from 0.025 to 0.35 molecules per amine group of the polyaminoamide.
 b) a water-soluble, crosslinked polyaminoamide derived by crosslinking a polyaminoamide as defined
above, with a crosslinking agent which is:

above, with a crosslinking agent which is:

10 I a bis-halogenohydrin bis-azetidinium compound, bis-halogenoacyl diamine or a bis-(alkyl halide),
Il an oligomer obtained by reacting a compound from group I, or an epihalogenohydrin, diepoxide or
bis-unsaturated derivative, with a difunctional compound or

Ill a product resulting from the quaternisation of a compound from group I or an oligomer from group II which contain one or more tertiary amine groups which can be alkylated with an alkylating agent, in an

amount from 0.025 to 0.35 molecule of crosslinking agent per amine group of the polyaminoamide, c) a water-soluble polyaminoamide derivative resulting from the condensation of a polyalkylenepolyamine with a polycarboxylic acid and alkylation of the condensate with a difunctional agent, (7) a polymer obtained by reacting a polyalkylenepolyamine, containing two primary amine groups and at

least one secondary amine group, with a dicarboxylic acid which is diglycolic acid or a saturated aliphatic dicarboxylic acid having 3 to 8 carbon atoms, the molar ratio of the polyalkylene-polyamine to the dicarboxylic acid being from 0.8:1 and 1.4:1 and the resulting polyamide being reacted with epichlorohydrin in a molar ratio of epichlorohydrin to the secondary amine group of the polyamide of 0.5:1 to 1.8:1.

(8) an optionally alkylated or alkoxylated polyalkylene-imine.(9) a product resulting from the condensation of a polyamine with epichlorohydrin.

25 (10) a quaternary polyureylene.
13. A composition according to any one of claims 1 to 12 in which the non-ionic surface-active agent is condensation product of a monoalcohol, an alpha-diol, an alkylphenol or an alkylamide with glycidol; a polyoxyethyleneated alcohol or alkylphenol, or an ester of a polyethylene glycol or polyglycerol with a linear fatty chain having 8 to 18 carbon atoms; a copolymer of ethylene oxide and propylene oxide, or a

fatty chain having 8 to 18 carbon atoms; a copolymer of ethylene oxide and propylene oxide, or a condensation product of ethylene oxide and propylene oxide with a fatty alcohol; a polyoxyethyleneated fatty amine; or a polyoxyethyleneated fatty acid ester of sorbitol or of sucrose.

14. A composition according to claim 13, in which the non-ionic surface-active agent is a compound of the formula:

35 (1) R₄-CHOH-CH₂-O-(CH₂-CHOH-CH₂-O)_p-H, in which R₄ denotes an aliphatic, cycloaliphatic or arylaliphatic radical having 7 to 21 carbon atoms, which can contain an ether, thioether, or hydroxymethylene group, and p is from 1 to 10; or (2) R₅O(C₂H₃O(CH₂OH))_aH,

in which R_s denotes an akyl, alkenyl or alkylaryl radical and q is from 1 to 10 or

(3) R₆CONH-CH₂-CH₂-O-CH₂-CH₂-O-CH₂-CH₂-OH₂-CH₂-OH₂-H, in which R₆ denotes a linear or branched, saturated or unsaturated, aliphatic radical, having 8 to 30 carbon atoms, which can contain one or more hydroxyl groups, and r is from 1 to 5.

15. A composition according to any one of claims 1 to 12, in which the surface-active agent containing one or more carboxyl or carboxylate groups in addition to non-ionic groups is a polyglyceryl carboxylate or carboxylic acid of a polyglycol ether, of the formula Alk-(OCH₂-CH₂)_n-OCH₂CO₂H, in which Alk denotes a linear chain having 12 to 18 carbon atoms and n is from 5 to 15, or a salt thereof.

16. A composition according to any one of claims 1 to 15, which contains one or more of an organic solvent, thickener, dyestuff, perfume, preservative, natural product, sequestering agent, emulsifier, softener, synergistic agent or foam stabiliser.

17. A composition according to claim 1 substantially as described in any one of the Examples
18. Process for treating fibrous material, which comprises applying thereto a composition as claimed in

any one of claims 1 to 17.

19. Process according to claim 18 in which after the application water is added to the material to deposit

19. Process according to claim 18 in which after the application water is added to the material to deposit the polymer onto the material.

20. Process according to claim 18 for treating human hair in which the composition is in the form of a shampoo, a colouring product, a rinsing lotion intended to be applied before or after shampooing, before or after colouring or bleaching or before or after permanent waving, a brushing lotion or a restructuring lotion.

21. Process according to claim 18 substantially as described in any one of the Examples.